

Postnatal development of the amygdala: a stereological study in macaque monkeys.

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Abstract

Abnormal development of the amygdala has been linked to several neurodevelopmental disorders, including schizophrenia and autism. However, the postnatal development of the amygdala is not easily explored at the cellular level in humans. Here, we performed a stereological analysis of the macaque monkey amygdala in order to characterize the cellular changes underlying its normal structural development in primates. The lateral, basal and accessory basal nuclei exhibited the same developmental pattern, with a large increase in volume between birth and three months of age, followed by slower growth continuing beyond one year of age. In contrast, the medial nucleus was near adult size at birth. At birth, the volume of the central nucleus was half of the adult value; this nucleus exhibited significant growth even after one year of age. Neither neuronal soma size, nor neuron or astrocyte numbers changed during postnatal development. In contrast, oligodendrocyte numbers increased substantially, in parallel with an increase in amygdala volume, after three months of age. At birth, the paralaminar nucleus contained a large pool of immature neurons that gradually developed into mature neurons, leading to a late increase in the volume of this nucleus. Our findings revealed that distinct amygdala nuclei exhibit different developmental profiles and that the amygdala is not fully mature for some time postnatally. We identified different periods during which pathogenic factors might lead to the abnormal development of distinct amygdala circuits, which may contribute to different human neurodevelopmental disorders associated with alterations of amygdala structure and functions.

Introduction

Abnormal structure or function of the amygdala has been implicated in the pathophysiology of a surprising number of psychiatric disorders including anxiety (Kalin et al., 2004; Kim et al., 2011; Tye et al., 2011), depression (Drevets, 2003; Hamidi et al., 2004; Yang et al., 2010), bipolar disorder (Berretta et al., 2007; Kalmar et al., 2009; Usher et al., 2010), schizophrenia (Kreczmanski et al., 2007; Lawrie et al., 2003) and autism (Baron-Cohen et al., 2000; Mosconi et al., 2009; Schumann and Amaral, 2006; Schumann et al., 2009; Schumann et al., 2004), as well as in neurological disorders such as epilepsy (Aliashkevich et al., 2003; Aroniadou-Anderjaska et al., 2008) and Alzheimer's disease (Cavedo et al., 2011; Ichinohe et al., 2009; Vereecken et al., 1994). To reconcile the diversity of disorders with which the amygdala is associated, one might speculate that there are sensitive developmental periods during which different environmental factors, hormonal influences or other experiences could alter specific aspects of amygdala development, and thus contribute to the etiologies of such a variety of psychiatric disorders (Tottenham and Sheridan, 2009). It is important, therefore, to understand the normal time course and features of amygdala development. However, the poor availability of early postnatal brain tissue makes a study such as this impractical in the human. Use of the nonhuman primate, therefore, is a valuable proxy for establishing the cellular processes underlying primate amygdala development.

To date, however, there is no reliable description of the morphological characteristics, such as cell numbers and volumes of the main amygdala nuclei, during the first years of life in primates. Previous studies have described the prenatal development of the monkey amygdala (Kordower et al., 1992; Ulfing et al., 2003). Their findings are consistent with work carried out in the human (Humphrey, 1968), indicating that the basic architecture of the amygdala is well established at birth. In contrast, MRI studies of postnatal amygdala development performed in humans and non-human primates indicate a delayed maturation of this structure that potentially extends up to the end of the second decade in humans (Giedd et al., 1996; Mosconi et al., 2009; Ostby et al., 2009; Payne et al., 2010; Schumann et al., 2004). Of course, current MRI techniques do not have sufficient resolution to quantify the volumetric changes of distinct amygdala nuclei nor to characterize, at the cellular level, the underlying morphological changes.

The goal of the present study was to provide quantitative neuroanatomical information about the postnatal maturation of the primate amygdala. We implemented design-based stereological techniques to characterize the cellular development of the rhesus monkey (*Macaca mulatta*) amygdaloid complex from birth to one year of age and in young adulthood (5-9 years of age). We aimed to determine when the volumes and cell numbers of its six main nuclei (i.e., lateral, basal, accessory basal, paralaminar, central and medial) reach an adult-like state. Understanding the normal time course of postnatal development of the primate amygdala, at the cellular level, will help to determine when specific structural and functional features develop, in order to be able to predict how these maturational processes can be affected by factors that might underlie the development of pathology in humans.

Materials and methods

Experimental animals

Twenty-four rhesus monkeys, *Macaca mulatta*; four 1-day-olds (2 males, 2 females), four 3-month-olds (2 M, 2 F), four 6-month-olds (2 M, 2 F), four 9-month-olds (2 M, 2 F), four 1-year-olds (2 M, 2 F) and four adults [5.3, 9.4 (M), 7.7 and 9.3 (F) years old] were used for this study. Monkeys were born from multiparous mothers and raised at the California National Primate Research Center (CNPRC). They were maternally reared in 2,000 m² outdoor enclosures and lived in large social groups until they were killed. These monkeys were the same animals used in quantitative studies of the monkey hippocampal formation (Jabès et al., 2010; 2011) and in a comparative study of the amygdala in young adult rats and monkeys (5-9-year-old monkeys only (Chareyron et al., 2011)). All experimental procedures were approved by the Institutional Animal Care and Use Committee of the University of California, Davis, and were conducted in accordance with the National Institutes of Health guidelines for the use of animals in research.

Brain acquisition

Monkeys were deeply anesthetized with an intravenous injection of sodium pentobarbital (50 mg/kg i.v., Fatal-Plus, Vortech Pharmaceuticals, Dearborn, MI) and perfused transcardially with 1% and then 4% paraformaldehyde in 0.1 M phosphate buffer (PB; pH 7.4) following protocols previously described (Lavenex et al., 2009).

Coronal sections were cut using a freezing, sliding microtome (Microm HM 450, Microm International, Germany) in six series at 30 μm and one series at 60 μm . The 60 μm sections were collected in 10% formaldehyde solution in 0.1 M PB (pH 7.4) and postfixed at 4°C for 4 weeks prior to Nissl staining with thionin. All other series were collected in tissue collection solution and kept at -70°C until further processing (Lavenex et al., 2009).

Histological processing

The procedure for Nissl-stained sections followed our standard laboratory protocol described previously (Chareyron et al., 2011; Lavenex et al., 2009). Briefly, sections were taken from the 10% formaldehyde solution, thoroughly washed, mounted on gelatin-coated slides, and air-dried overnight at 37°C. Sections were then defatted 2 X 2 hours in a mixture of chloroform/ethanol (1:1, vol.), partially rehydrated and air-dried overnight at 37°C. Sections were then fully rehydrated and stained 20 seconds in a 0.25% thionin solution (Fisher Scientific, Waltham, MA, cat# T-409), dehydrated and coverslipped with DPX (BDH Laboratories, Poole, UK).

Anatomical boundaries of the amygdala

The nomenclature and the basic description of the morphological characteristics of the amygdala nuclei have been described in detail previously by Price et al. (1987), Amaral et al. (1992) and Pitkänen and Amaral (1998) for the cynomolgus monkey (*Macaca fascicularis*). We used these descriptions to determine the boundaries of the six main nuclei (i.e., lateral, basal, accessory basal, paralaminar, central and medial) of the *Macaca mulatta* amygdala (Fig. 1) (Chareyron et al., 2011).

Stereological analyses

Volume measurements, and neuron and glia counts were performed with Stereoinvestigator 9.0 (Microbrightfield, Williston, VT). We estimated the volume of the brain (telencephalon and diencephalon), the volume of the whole amygdala, the volume of the main amygdala nuclei (lateral, basal, paralaminar, accessory basal, central and medial) according to the Cavalieri principle on Nissl-stained sections cut at 60 μm (Gundersen and Jensen, 1987; Lavenex et al., 2000a; b; West and Gundersen, 1990). We used the section cutting thickness (60 μm) to calculate the volume. Brain volume refers to the volume of the telencephalon and diencephalon

bilaterally (ventricles were excluded). Twenty-nine to thirty-seven sections per animal (1,920 μm apart), with the first section selected randomly within the first seven sections through the brain, were used for brain volume measurements. An average of fourteen sections per animal (480 μm apart), with the first section selected randomly within the first two sections through the amygdala, was used to measure the volume of the whole amygdala (for a complete list of amygdala nuclei, see (Chareyron et al., 2011)). We compared total amygdala volumes in the left and the right hemisphere and did not find evidence of lateralization ($F_{1,23} = 0.0977$, $P = 0.757$). We therefore performed estimates unilaterally for the other parameters (see below). About twenty-eight sections per animal (240 μm apart) were used for volume estimates of the main amygdala nuclei. At the rostral pole of the amygdala, the paralaminar nucleus extends rostrally and dorsally to the lateral and basal nuclei. As a consequence, the first coronal section may be oriented tangentially to the paralaminar nucleus, resulting in an over-representation of this region in some cases. We therefore excluded this rostral, dorsal portion of the paralaminar nucleus, and restricted our analysis to the main part of the nucleus located ventrally to the basal nucleus. Since there was no lateralization for overall amygdala size, we estimated the volumes of individual amygdala nuclei in the left hemisphere for half of the animals, and in the right hemisphere for the other half (balanced across sexes).

The total numbers of neurons in the main amygdala nuclei were estimated using the optical fractionator method (Gundersen, 1986; West et al., 1991). This design-based method enables an estimation of cell number that is independent of volume estimates. Neuron number was estimated in the right or in the left amygdala only, as for volume measurements. About nine sections per animal (480 μm apart), with the first section selected randomly within the first two sections through the nucleus of interest, were used for neuron counts (Table 1). We used a 100X PlanFluor oil objective (N.A. 1.30) on a Nikon Eclipse 80i microscope (Nikon Instruments Inc, Melville, NY, USA) linked to PC-based StereoInvestigator 9.0. The sampling scheme was established to obtain individual estimates of neuron number with estimated coefficients of error (CE) around 0.10 (CE average (neurons) = 0.121). This sampling scheme was the same as in our previous study of the adult monkey amygdala (Chareyron et al., 2011), except for the basal nucleus. Section thickness was measured at every other counting site (Table 1).

The volumes of neuronal somas were determined using the Nucleator method (Gundersen, 1988). We measured an average of two hundred and sixty neurons per nucleus, sampled at every counting site during the optical fractionator analysis. Briefly, the nucleator can be used to estimate the mean cross-sectional area and volume of cells. A set of rays emanating from a point randomly chosen within the nucleus is drawn and oriented randomly. The length of the intercept from the point to the cell boundary (l) is measured and the cell volume is obtained by $V = (4/3) \times 3.1416 \times (\text{mean } l)^3$. Essentially, this is the formula used to determine the volume of a sphere with a known radius. Note that the nucleator method provides accurate estimates of neuron size when isotropic-uniform random sectioning of brain structures is employed (Gundersen, 1988). In our study, all brains were cut in the coronal plane. Therefore, although estimates of cell size might be impacted by the non-random orientation of neurons in the different amygdala nuclei, which could lead to a systematic over- or under-estimation of cell size in any given structure, comparisons between cell sizes in the same structure across developmental ages should not be impacted.

The total number of glial cells in the main amygdala nuclei was estimated using the optical fractionator method during neuron counting. Thus, the same sampling scheme used for neuron counts was used for glial cell counts (CE average (astrocytes) = 0.115, CE average (oligodendrocytes) = 0.275). We distinguished neurons, oligodendrocytes and astrocytes based on morphological criteria identifiable in Nissl preparations (Chareyron et al., 2011) (Fig. 2A,B,C). We refer the reader to the original publications by (Fitting et al., 2008; Grady et al., 2003; Hamidi et al., 2004; Palackal et al., 1993) for detailed descriptions. Briefly, neurons are darkly stained and comprise a single large nucleolus. Astrocytes are relatively smaller in size and exhibit pale staining of the nucleus. Oligodendrocytes are smaller than astrocytes and contain round, darkly-staining nuclei that are densely packed with chromatin. Microglia were not counted but could be identified because they have the smallest nucleus, dark staining, and an irregular shape that is often rod-like, oval or bent (Morris et al., 2008). We also identified a population of immature cells in the paralaminar nucleus (Fig. 2D). These cells were small with round to slightly oval, hyperchromatic nuclei containing distinguishable nucleoli (Bernier et al., 2002; Fudge, 2004; Yachnis et al., 2000).

Photomicrographic production

Low-magnification photomicrographs were taken with a Leica DFC420 digital camera on a Leica MZ9.5 stereomicroscope (Leica Microsystems, Wetzlar, Germany). High-magnification photomicrographs were taken with a Leica DFC490 digital camera on a Nikon Eclipse 80i microscope (Nikon Instruments, Tokyo, Japan). Artifacts located outside of the sections were removed, and levels were adjusted in Adobe Photoshop CS4, v. 11.0 (Adobe, San Jose, CA) to improve contrast and clarity.

Statistics

We performed analyses of variance (ANOVAs) with age as a factor on the estimates of the volume, mature neuron number, immature neuron number, astrocyte and oligodendrocyte numbers and neuronal soma size, as these data were normally distributed. Post hoc analyses were performed with the Fisher-PLSD test. Significance level was set at $P < 0.05$ for all analyses. No consistent sex difference was found for the estimated parameters (data not shown), so data from both sexes were combined for presentation. We also evaluated both left and right amygdala nuclei in a systematic manner (as described above) and no consistent lateralization was found (data not shown). Thus, our findings and all subsequent considerations are valid for, and can be generalized to, both left and right amygdala in both males and females. All sections used in this study were coded to allow blind analysis, and the code was broken only after completion of the analyses (with the exception of the sections from the 5-9-year-old monkeys that were analyzed first and published in a separate article (Chareyron et al., 2011)). All analyses were performed by the same experimenter, LJC.

Results

Volumes

The volumes of the monkey brain, amygdala and its six main nuclei, at different postnatal ages, are summarized in Table 2. Note that all these volumes were estimated, following brain fixation by perfusion with 4% paraformaldehyde, using frozen coronal sections cut at 60 μm and Nissl-stained with thionin. Due to fixation and processing-induced shrinkage, the resulting volumes are substantially less than the estimates of in vivo brain volumes obtained with MRI technology (see below for

discussion). The volume of the brain (telencephalon and diencephalon, bilaterally) differed between age groups ($F_{5,18} = 3.254$, $P = 0.029$). At birth, brain volume was about 85% of the adult volume, which was reached by three months of age (newborn < all other ages, all $P < 0.044$). The volume of the whole amygdala also differed between age groups ($F_{5,18} = 9.308$, $P < 0.001$). At birth, amygdala volume was only 60% of the adult volume; it was significantly larger at three months of age reaching about 82% of the adult volume (newborn < 3-month-olds < adults, all $P < 0.006$).

The volumes of the lateral, basal and accessory basal nuclei exhibited similar developmental profiles, whereas those of the paralamina, central and medial nuclei demonstrated differences (Table 2). At birth, the volume of the lateral nucleus was 66% of its adult value (Fig. 3A); it was about 91% of the adult value at three months of age ($F_{5,18} = 5.025$, $P = 0.005$; newborn < 3-month-olds, $P = 0.003$). At birth, the volume of the basal nucleus was 55% of the adult volume (Fig. 3A); it was significantly larger at three months of age reaching about 72% of the adult volume ($F_{5,18} = 7.212$, $P < 0.001$; newborn < 3-month-olds < adults, all $P < 0.035$). Similarly, the volume of the accessory basal nucleus was 57% of its adult volume at birth (Fig. 3A) and was significantly larger at three months of age reaching about 78% of the adult volume ($F_{5,18} = 7.697$, $P < 0.001$; newborn < 3-month-olds < adults, all $P < 0.009$).

In contrast, the paralamina nucleus was only at 30% of its adult size at birth (Fig. 3B; $F_{5,18} = 16.144$, $P < 0.001$); it differed between birth and six months of age (when it reached 50% of adult size; $P = 0.035$), as well as between birth and one year (55% of adult size) and 5-9-years of age ($P < 0.001$). The volume of the central nucleus was only 50% of its adult size at birth (Fig. 3B; $F_{5,18} = 15.031$, $P < 0.001$). At one year of age, it reached only 57% of its adult volume (1-year-olds < 5-9-year-olds, $P < 0.001$). Finally, although the volume of the medial nucleus increased from 85% of its adult value at birth, the differences between ages were not significant (Fig. 3B; $F_{5,18} = 1.807$, $P = 0.162$).

Neuron numbers

There were no differences in the numbers of neurons estimated at different postnatal ages in five of the main amygdala nuclei in monkeys (Table 2): lateral nucleus ($F_{5,18} = 2.240$, $P = 0.095$), basal nucleus ($F_{5,18} = 1.178$, $P = 0.358$),

accessory basal nucleus ($F_{5,18} = 0.543$, $P = 0.742$), central nucleus ($F_{5,18} = 0.791$, $P = 0.570$) and medial nucleus ($F_{5,18} = 1.339$, $P = 0.293$).

In contrast, the paralamina nucleus contained both mature neurons and a pool of morphologically distinct, immature cells (Fig. 2D). The number of mature neurons in this nucleus differed between age groups ($F_{5,18} = 92.284$, $P < 0.001$; Table 2). From birth to one year of age, the nucleus contained only 15% of the total number of mature neurons observed in 5-9-year-old monkeys (all other ages < 5 -9-year-olds, all $P < 0.001$; Fig. 4A). In order to verify that these differences were not due to differences in the delineation of the nucleus at different ages, especially at its border with the basal nucleus, we evaluated the differences in neuron numbers summed across both nuclei i.e., the sum of the neurons in the paralamina and basal nuclei. We found that when the two nuclei were grouped together, there were similar differences between ages ($F_{5,18} = 7.364$, $P < 0.001$; all other ages < 5 -9-year-olds, all $P < 0.001$) as when the paralamina nucleus was evaluated alone; as noted above, the neuron number in the basal nucleus alone did not differ in this way. The increase of about 350,000 mature neurons in the paralamina nucleus was paralleled by the disappearance of an equivalent number of morphologically distinct, presumably immature neurons ($F_{5,18} = 3.965$, $P = 0.013$; all other ages > 5 -9-year-olds, all $P < 0.023$; Table 2; Fig. 4A). It is important to note that the sum of the number of mature neurons and immature cells observed in the paralamina nucleus did not differ between ages ($F_{5,18} = 0.624$, $P = 0.684$). This supports the idea that these morphologically distinct cells are immature neurons that are present in the paralamina nucleus at birth but mature after one year of age.

Astrocyte numbers

There were no significant differences in the numbers of astrocytes found in five of the main amygdala nuclei of monkeys at different postnatal ages (Table 2): lateral nucleus ($F_{5,18} = 2.562$, $P = 0.064$), basal nucleus ($F_{5,18} = 0.526$, $P = 0.754$), accessory basal nucleus ($F_{5,18} = 2.251$, $P = 0.093$), central nucleus ($F_{5,18} = 0.720$, $P = 0.617$) and medial nucleus ($F_{5,18} = 1.011$, $P = 0.440$).

In contrast, the number of astrocytes in the paralamina nucleus differed between age groups ($F_{5,18} = 5.308$, $P = 0.004$; Fig. 4); it was lower in newborn, as compared to 3-month-old monkeys ($P = 0.01$). The numbers in monkeys at the time points

between 3 months and 1 year of age, were lower than in 5-9-year-old monkeys (3-month, 6-month, 9-month, 1-year < 5-9-year; all $P < 0.007$).

Oligodendrocyte numbers

The numbers of oligodendrocytes differed between age groups in the six main amygdala nuclei (Table 2; Fig. 5). For the lateral nucleus (Fig. 5A; $F_{5,18} = 46.777$, $P < 0.001$), newborns had fewer cells than 9-month-olds and older (all $P < 0.047$) and 1-year-olds had fewer cells than 5-9-year-olds ($P < 0.001$). In the basal nucleus (Fig. 5B; $F_{5,18} = 95.712$, $P < 0.001$), newborns had fewer cells than 6-month-olds, which had less than 1-year-olds, which again had less than 5-9-year-olds (all $P < 0.026$). For the accessory basal nucleus (Fig. 5C; $F_{5,18} = 77.443$, $P < 0.001$), newborns had fewer cells than 9-month-olds and older (all $P < 0.008$) and 1-year-olds had fewer oligodendrocytes than 5-9-year-olds ($P < 0.001$). In the paralamina nucleus (Fig. 5D; $F_{5,18} = 37.087$, $P < 0.001$), all younger ages had fewer oligodendrocytes than 5-9-year-olds (all $P < 0.001$). For the central nucleus (Fig. 5E; $F_{5,18} = 40.163$, $P < 0.001$), newborns had fewer oligodendrocytes than 6-month-olds, which had fewer cells than 1-year-olds, which had lower numbers than 5-9-year-olds (all $P < 0.039$). Finally, in the medial nucleus (Fig. 5F; $F_{5,18} = 17.524$, $P < 0.001$), newborns had fewer cells than 6-month-olds ($P = 0.014$) and 1-year-olds had fewer cells than 5-9-year-olds ($P < 0.001$).

Neuron soma size

We did not find any age-related differences in average neuronal soma size in five of the main amygdala nuclei (Table 2; Fig. 6): lateral nucleus ($F_{5,18} = 1.052$, $P = 0.418$), basal nucleus ($F_{5,18} = 1.468$, $P = 0.249$), accessory basal nucleus ($F_{5,18} = 0.631$, $P = 0.679$), central nucleus ($F_{5,18} = 1.822$, $P = 0.159$) and medial nucleus ($F_{5,18} = 1.309$, $P = 0.304$).

In contrast, average neuronal soma size differed between age groups in the paralamina nucleus (Fig. 7; $F_{5,18} = 4.176$, $P = 0.011$); newborns had smaller neurons than 6-month-olds and 1-year-olds (all $P < 0.008$) and 3-month-olds had smaller neurons than 5-9-year-olds ($P = 0.044$). The percentage of paralamina neurons with a soma volume above $1,500 \mu\text{m}^3$ (the average adult neuronal soma volume) differed between ages within the first year of postnatal life ($F_{4,15} = 4.921$, $P = 0.01$). In newborns, this percentage was only 6.0%; it increased significantly ($P = 0.004$) to

38.8% in 1-year-olds. These differences were linked to the maturation of neurons already present in the paralaminar nucleus at birth (Fig. 4; as described above).

Volume and oligodendrocyte numbers

During the first three months of postnatal life, amygdala nuclei exhibited increases in volume that were not related to any changes in the cellular processes evaluated in the current study. This indicates that the increased volume was due to an increase in neuropil volume. Between three months and 5-9 years of age, increases in the volumes of individual amygdala nuclei were not correlated with the numbers of neurons or astrocytes nor with the volume of neuronal soma. In contrast, after three months of age, the volume of the whole amygdala correlated with the total number of oligodendrocytes (volume = $96.627 + 0.0000108 \times \text{oligodendrocyte number}$; $R^2=0.58$; $F_{1,18} = 25.349$; $P < 0.001$). A significant linear relationship between volume and oligodendrocyte number was observed for all nuclei, except for the medial nucleus: lateral nucleus (volume = $32.747 + 0.0000053 \times \text{oligodendrocyte number}$; $R^2=0.23$; $F_{1,18} = 5.636$; $P = 0.029$; Fig. 8A), basal nucleus (volume = $33.679 + 0.000011 \times \text{oligodendrocyte number}$; $R^2=0.49$; $F_{1,18} = 17.523$; $P < 0.001$; Fig. 8B), accessory basal nucleus (volume = $19.259 + 0.0000092 \times \text{oligodendrocyte number}$; $R^2=0.42$; $F_{1,18} = 13.129$; $P = 0.002$; Fig. 8C), paralaminar nucleus (volume = $3.782 + 0.0000482 \times \text{oligodendrocyte number}$; $R^2=0.65$; $F_{1,18} = 34.620$; $P < 0.001$; Fig. 8D), central nucleus (volume = $2.904 + 0.0000232 \times \text{oligodendrocyte number}$; $R^2=0.81$; $F_{1,18} = 76.982$; $P < 0.001$; Fig. 8E), medial nucleus (volume = $4.548 + 0.00000686 \times \text{oligodendrocyte number}$; $R^2=0.18$; $F_{1,18} = 4.169$; $P = 0.056$; Fig. 8F). Overall, these data suggest that increases in volume after three months of postnatal life are linked, at least in part, to increases in oligodendrocyte numbers and the myelination of fibers within the amygdala. Interestingly, this relation was strongest for the central nucleus. Other changes, including the maturation of dendritic arborization are also likely to contribute to postnatal volumetric changes. Analysis of postnatal dendritic maturation in several nuclei of the rhesus monkey is currently being undertaken.

Discussion

The goal of the present study was to provide quantitative neuroanatomical information about the cellular postnatal maturation of the rhesus monkey amygdala,

including when the volumes and cell numbers of its six main nuclei (i.e., lateral, basal, accessory basal, paralaminar, central and medial) reach an adult-like state. Our three major findings are as follows: (1) amygdala nuclei exhibited different developmental profiles; (2) neuronal soma size, and neuron and astrocyte numbers did not vary after birth in five of the six main amygdala nuclei, whereas the numbers of oligodendrocytes increased continuously in all nuclei from birth to beyond one year of age; (3) at birth, the paralaminar nucleus was comprised of a group of mature neurons and a group of immature neurons that developed after one year of age.

There is, to our knowledge, no published information on the postnatal maturation of the monkey or human amygdala at the cellular level. We therefore compare our results to non-invasive MRI studies carried out in primates, including monkeys and humans, as well as to previous work carried out at the cellular level in rats and rabbits. Our systematic, quantitative findings provide detailed information regarding the cellular changes underlying the different developmental profiles of distinct amygdala nuclei in monkeys, which are largely consistent with, and expand upon, the findings of previous reports in other species.

Technical considerations

First, it is important to consider that age-related, differential shrinkage of brain tissue during processing cannot explain the age differences in the volumes of distinct amygdala nuclei that we observed. Although the average thickness of processed sections was lower in newborn cases (9.6 μm or 16% of the cutting thickness), as compared to all other ages (13.0 μm or 22% of the cutting thickness), volume estimates were based on the cutting section interval (240 μm for individual amygdala nuclei). The impact of age-dependent differential shrinkage in the x- and y-plane of frozen brain sections is minimal and likely minor, as compared to shrinkage in the z-plane (Carlo and Stevens, 2011). In addition, our data derived from estimates made on frozen sections cut at 60 μm and Nissl-stained with thionin are consistent with the direction and magnitude of changes in the volume of the amygdala estimated in-vivo with non-invasive MRI technology (even though estimates based on histological preparations are consistently smaller (by about 40% in mature individuals) than in-vivo estimates (i.e. telencephalon+diencephalon in 5-9-year-old monkeys: 52,360 mm^3 (present study) vs. whole brain – (cerebellum+brainstem) in 8-12-year-old monkeys: 87,080 mm^3 (Scott et al., Submitted); see below for further discussion).

Moreover, since we observed different patterns of postnatal development for distinct amygdala nuclei it is unlikely that differential shrinkage impacts the suggestions regarding the relative development of specific circuits and processes that can be derived from our volumetric data. Importantly, cell counts are not impacted by differential shrinkage, as the optical fractionator technique provides estimates of cell numbers that are independent of volume estimates. Similarly, estimates of neuronal soma size are likely not significantly affected, as we did not observe any significant developmental increase (except for the paralaminar nucleus) in neuronal soma size.

Brain and amygdala volumes

The rapid increase in brain volume that we observed during the first three months of life in rhesus monkeys is consistent with previous findings of MRI studies in monkeys (Malkova et al., 2006; Payne et al., 2010; Scott et al., Submitted) and humans (Courchesne et al., 2000; Giedd et al., 1999; Knickmeyer et al., 2008; Pfefferbaum et al., 1994).

In rhesus monkeys, Malkova et al. (2006) reported that brain volume is 64% that of the young adult at one week of age. The most substantial increase occurs between one week and two months followed by lesser increases until three years of age. Similarly, Payne et al. (2010) reported that at one week of age, brain volume is about 60% that of the 2-year-old monkey. Most recently, Scott et al. (Submitted) found that the average brain volume of one week-old monkeys is 66% that of 9-12-year-olds. By three months of age, brain volume has reached 85% of its adult value, and by one year, it has reached adult volume. These MRI findings, which are largely consistent, differ from our brain volume estimates in perfusion-fixed tissue (Table 2). This difference might be due to the shrinkage that occurs during perfusion-fixation and freezing of the monkey brains, before they are cut and the sections are stained. In contrast to the shrinkage in the z axis produced during the processing of individual 60 μm sections, which is greater in newborns as compared to older ages (see above), surprisingly, overall brain shrinkage following perfusion and freezing processes appears to be greater in older monkeys (post-mortem brain volume is 61% of the in vivo volume in 5-9-year-olds) than in younger monkeys (post-mortem brain volume is 77% of the in vivo volume in newborns (Table 2). At this time, we are at a loss to explain such age-dependent differential shrinkage. This suggests that the developmental increases in volume of individual amygdala nuclei that we estimated

from post-mortem material might be even larger in live subjects. Importantly, the relatively greater (in relation to overall brain size) enlargement of the monkey amygdala that we observed between birth and three months of age is in agreement with reported data on amygdala development based on MRI analyses (Payne et al., 2010).

Our data in monkeys are also consistent with MRI studies performed in humans that show a linear increase in size of the amygdala of about 75 mm^3 per year from 18 to 59 months of age (Mosconi et al., 2009; Ostby et al., 2009), which continues into young adulthood at about 70 mm^3 per year (Giedd et al., 1996; Ostby et al., 2009; Schumann et al., 2004). To our knowledge, there are no data on the development of the human amygdala between birth and 2 years of age, which would enable a better comparison of our results with those obtained during the corresponding developmental period in humans.

Nuclei volumes & cell numbers

Our findings of differential postnatal development of distinct amygdala nuclei in monkeys are consistent with previously published data in rats and rabbits, despite the fact that, at birth, primates are comparatively more mature than both rats and rabbits. Similar to the early and rapid maturation of the lateral, basal and accessory basal amygdala nuclei that we observed in monkeys between birth and three months of age, the rat lateral and basal amygdala nuclei increase rapidly in size from birth to three weeks of postnatal age (P21), where their combined volume reaches about 90% of the volume observed at six months of age (10% at P0, 40% at P7, and 80% at P14; (Berdel et al., 1997b)). This period of rapid enlargement is followed by a much slower maturation that extends beyond three months of age, when the basolateral amygdala has reached about 95% of the volume observed in six-month-old rats (Berdel et al., 1997b; Rubinow and Juraska, 2009). In the rabbit, the combined volume of the lateral, basal and accessory basal nuclei increases linearly from birth (12% of adult value) to four months of age, when it reaches the value observed in six-month-old animals (Jagalska-Majewska et al., 2003). Interestingly, and similar to what we found in monkeys, the medial nucleus reaches an adult volume between P5 and P11 in rats (Mizukami et al., 1983), an age at which the combined volume of the lateral and basal nuclei is still less than 80% of the adult value (Berdel et al., 1997b). Although the actual percentages of adult values differ

between species, these data indicate that the medial nucleus matures relatively earlier than the lateral and basal nuclei in all species examined.

Considering that primates are born significantly more mature than rats and rabbits, the lack of postnatal changes in neuron numbers and neuronal soma volumes in monkeys are consistent with data in rats indicating a stable number of neurons after P7 and an adult-like neuronal soma size at P14 (Berdel et al., 1997a; Rubinow and Juraska, 2009). Nevertheless, a systematic stereological study of the postnatal development of the rat amygdala will be necessary to make direct comparisons and draw more meaningful parallels regarding the similarities or differences in the cellular maturational processes among different species.

Previous studies reported that postnatal neurogenesis occurs in the amygdala of mice (Shapiro et al., 2009), voles (Fowler et al., 2002) and monkeys (Bernier et al., 2002). In contrast, others have shown that although an enriched environment increases the proliferation of cells in the mouse amygdala, almost all of the cells express the oligodendrocyte progenitor marker Olig2 (Okuda et al., 2009). Our systematic quantitative analysis, using design-based stereological techniques, failed to detect any changes in mature neuron numbers in five of the main monkey amygdala nuclei from birth to adulthood. Interestingly, postnatal neurogenesis is much more prominent and unambiguously demonstrated in the dentate gyrus, but even in this brain region, neurogenesis does not lead to changes in the total number of neurons in adult monkeys (that is, after a defined, postnatal developmental period (Jabès et al., 2010)). In contrast, we found an increase in mature neuron number in the paralaminar nucleus from one year to 5-9 years of age, which appears to be due to the postnatal maturation of immature neurons already present at birth. The presence of immature neurons in the paralaminar nucleus is consistent with the presence of Bcl-2-positive cells reported previously (Bernier et al., 2002).

In sum, our quantitative estimates indicate that whether or not marginal postnatal neurogenesis exists in the monkey amygdala, it does not lead to detectable postnatal changes in total neuron numbers.

Differential maturation of distinct amygdala circuits

Our stereological data identified different patterns and stages of postnatal development that might reflect the maturation of distinct cellular processes and amygdala circuits (Fig. 9). The first stage was characterized by the dramatic

enlargement of the deep nuclei of the amygdala (lateral, basal and accessory basal nuclei) between birth and three months of age. The second stage was characterized by a continuous increase in size of all amygdala nuclei beyond one year of age. In addition, the central and paralamina nuclei exhibited a notably slow and prolonged maturation.

First stage: neuropil expansion. In the absence of detectable changes in neuronal soma size or cell numbers during the first three months of life, the dramatic enlargement of the lateral, basal and accessory basal nuclei of the monkey amygdala likely reflects an increase in neuropil volume. Combined, these three nuclei exhibited a 35% increase in volume between birth and three months of age. Interestingly, these three nuclei represent about 56% of the volume of the entire amygdala in monkeys, and the volumetric changes observed for the whole amygdala exhibited the same developmental profile as that observed for these nuclei. Volume estimates based on MRI analyses in monkeys are therefore likely to be dominated by the developmental changes in these three nuclei, whereas more subtle changes in other amygdala nuclei are likely to be masked. This is probably even more the case in humans where these three nuclei represent about 69% of the size of the entire amygdala (Chareyron et al., 2011). Importantly, we did not find any changes in the numbers of neurons, astrocytes or oligodendrocytes, or in the size of neuronal somas in these three nuclei during the first three months of postnatal life. It is thus reasonable to assume that neuropil expansion underlies the observed volumetric changes. Indeed, the neuropil, which is defined as the portion of neural tissue composed mostly of small dendrites and spines and fine glial processes, unmyelinated axons and axon terminals (Bourgeois et al., 1994; Bourgeois and Rakic, 1993), represents the largest fraction of these nuclei. In contrast, the average volume occupied by neuronal somas in these three nuclei decreases from 11.2% at birth to 9.2% at three months of age and eventually reaches 6.3% at 5-9 years of age.

An increase of neuropil volume might, in turn, reflect an increase in the connectivity of these nuclei. The lateral, basal and accessory basal nuclei are highly interconnected with the neocortex (Amaral et al., 1992) and their volumetric expansion during the first three months of life parallels the volumetric expansion of the rest of the brain (current study; Malkova et al., 2006; Payne et al., 2010; Scott et al., Submitted). Interestingly, the massive volumetric expansion of prefrontal cortical

areas in the monkey has been linked to neuropil expansion during the first two months of postnatal life (Bourgeois et al., 1994). In rats, major changes in the interconnections between the basolateral amygdala and the prefrontal cortex occur between P7 and P19 (Bouwmeester et al., 2002a; Bouwmeester et al., 2002b). In cats, an increase in total dendritic length in the basolateral amygdala has been observed during the first postnatal month (Wakefield and Levine, 1985). These observations suggest that neuropil expansion related to changes in interconnectivity with neocortical areas might underlie the volumetric increases of the lateral, basal and accessory basal nuclei during the first three months of life. In contrast to the deep nuclei, the central and medial nuclei do not exhibit any increase in volume between birth and three months of age. These two nuclei are connected mainly with the thalamus, hypothalamus and brainstem (Amaral et al., 1992). In contrast to cortical connections, the interconnections between the amygdala and the thalamic regions and substantia innominata do not change between P7 and P19 in rats, suggesting that these connections are already established shortly after birth (Bouwmeester et al., 2002a; Bouwmeester et al., 2002b). The lack of significant volumetric changes of the monkey central and medial nuclei during the first postnatal year might thus reflect the lack of structural (and potentially functional) refinement of the interconnections between these nuclei and the rest of the brain. However, what is puzzling is the fact that the central nucleus exhibits a very significant enlargement after one year of age.

Second stage: myelination. We observed a continuous increase in the size of all amygdala nuclei from birth to young adulthood that was associated with an increase in the number of oligodendrocytes. This suggests that myelination of the amygdala circuits may underlie, at least in part, the late and continuous enlargement of amygdala nuclei after three months of age. Indeed, oligodendrocytes produce the myelin sheath that surrounds axons (Nave, 2010), and an increase in oligodendrocyte number has been linked to an increase in axon myelination during development (O'Kusky and Colonnier, 1982). The contribution of oligodendrocyte processes to the volumetric changes observed after three months of age was, however, not uniform across distinct amygdala nuclei. In the lateral, basal and accessory basal nuclei, statistical analyses indicated that oligodendrocyte numbers were only partially related to the volumetric changes. Thus, other factors might also contribute to these changes. In the medial nucleus, the correlation was also low due

to (a) high biological variability in this nucleus, and (b) a possible decrease in neuropil volume that has been suggested to occur during puberty (Zehr et al., 2006). In contrast, the correlation between volume and oligodendrocyte number was very high in the central nucleus, suggesting that myelination of fibers in this nucleus might contribute most to its volumetric maturation. In sum, despite subtle differences between individual amygdala nuclei, increases in oligodendrocyte number might underlie the continuous increase in overall amygdala volume until young adulthood.

Paralaminar nucleus

We quantified a population of immature cells in the paralaminar nucleus of the postnatal monkey amygdala, which developed into mature neurons after one year of age. Previous studies have noted the existence of immature cells expressing different markers of immature neurons in the paralaminar nucleus of: (1) monkeys: doublecortin (Decampo and Fudge, 2011; Marlatt et al., 2011; Zhang et al., 2009), PSA-NCAM (Bernier et al., 2002; Decampo and Fudge, 2011; Marlatt et al., 2011), Bcl2 protein (Bernier et al., 2002; Bernier and Parent, 1998a; b; Fudge, 2004), class III β -tubulin and TUC-4 (Bernier et al., 2002); (2) humans: Bcl-2 protein (Yachnis et al., 2000); and (3) rabbits: class III β -tubulin and PSA-NCAM (Luzzati et al., 2009; Luzzati et al., 2003), doublecortin and Tbr-1 (Luzzati et al., 2009). In particular, Yachnis et al. (2000) identified a population of immature cells in the ventrolateral region of the human amygdala, which are abundant throughout infancy, persist into adulthood, and are gradually reduced during adult life. Zhang et al. (2009) also reported that, although the number of doublecortin-expressing cells in the paralaminar nucleus decreases in adulthood, they remained present in 12-, 21- and even 31-year-old monkeys, suggesting continued neuronal maturation throughout the lifespan. Thus, although several studies have previously provided evidence of the presence of immature cells in the paralaminar nucleus, our study is the first to quantify this population of immature cells and their development during early postnatal life. The postnatal maturation of this population of neurons should not be confounded with the temporal stream reported by Bernier and colleagues (Bernier et al., 2002). Although we cannot exclude the possibility that some immature neurons migrate, the fact that we found that their number does not change during postnatal development suggests that these cells were quiescent in the paralaminar nucleus and only matured after one year of age.

What needs to be investigated now is whether these late maturing neurons in the paralamina nucleus integrate into existing amygdala circuits and, if so, what their function might be. The delayed maturation of these neurons must constitute a prominent phenomenon as these neurons represented 8% of the total number of mature neurons present in the basolateral portion of the amygdala (i.e., including the lateral, basal, accessory basal and paralamina nuclei) in young adult individuals. Unfortunately, only scant information is available about the connectivity of the paralamina nucleus and there is no clear hypothesis about its function (Amaral et al., 1992; Decampo and Fudge, 2011). Some authors assume that the integration of new (i.e., recently mature) neurons in the amygdaloid complex may parallel the continuing addition of neurons in the olfactory bulb, as the amygdala receives direct olfactory inputs (Arisi et al., 2011; Bernier et al., 2002). This hypothesis is problematic, however, as the paralamina nucleus, defined by the presence of immature cells, is undetectable in the rat (Chareyron et al., 2011). The amygdala is also interconnected with the hippocampus (Amaral et al., 1992; Decampo and Fudge, 2011) and as observed in the dentate gyrus (Jabès et al., 2010), the formation of new emotional memories could be facilitated by the integration of new neurons in the amygdala. Although a clear hypothesis about the function of the paralamina nucleus is still lacking, the addition of recently mature neurons to existing circuits in adolescent and adult animals might contribute to increased plasticity and coding capacity in the amygdala (Decampo and Fudge, 2011).

Conclusion

We identified two stages of postnatal development of the primate amygdala. The first is characterized by a large increase in the volumes of the lateral, basal and accessory basal nuclei between birth and three months of age, without any detectable changes in neuronal soma size or cell numbers, thus suggesting an increase in neuropil volume. The second stage is characterized by a slow and continuous increase in size of most amygdala nuclei from birth to young adulthood, which is accompanied by an increase in oligodendrocyte number. This stage likely reflects the postnatal myelination of amygdala circuits. We also quantified the maturation of a population of immature cells present in the paralamina nucleus at birth, which further develop into mature neurons after one year of age. Our quantitative findings in monkeys have defined the normal time course of postnatal

development of the primate amygdala at the cellular level. These data, which are likely a reasonable proxy for the development of the human amygdala, may provide the substrate for the formulation of models relating the perturbation of specific maturational processes to the etiologies of human psychiatric and neurological disorders.

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FIGURE LEGENDS

Figure 1: Low-magnification photomicrographs of a representative coronal section at a mid-rostrocaudal level of the rhesus monkey (*Macaca mulatta*) amygdala, which illustrate the locations of the six main amygdala nuclei in a newborn (**A** and **B**) and a 9-year-old (**C** and **D**) monkey. L: Lateral; B: Basal; PL: Paralaminar; AB: Accessory basal; CE: Central; M: Medial. Non-labeled areas include the remaining nuclei of the amygdala (see (Chareyron et al., 2011)). Scale bar in **A**: 1 mm, applies to all panels.

Figure 2: Classification and identification of different cell types in the basal (A,B,C) and paralaminar (D) nuclei of the monkey amygdala, viewed with an X100 objective in Nissl-stained, coronal sections cut at 60 μm . **A**: Neuron. **B**: Astrocytes. **C**: Oligodendrocytes. **D**: Immature neurons. Scale bar in **A**: 5 μm , applies to all panels.

Figure 3: Volumes of the main nuclei of the monkey amygdala at different ages during early postnatal development (expressed as a percentage of the volume of the structure observed in 5-9-year-old monkeys: average \pm SD). **A**: Lateral, basal and accessory basal nuclei. **B**: Paralaminar, central and medial nuclei. N: Newborn; 3M: 3-month-olds; 6M: 6-month-olds; 9M: 9-month-olds; 1Y: 1-year-olds; 5-9Y: 5-9-year-olds.

Figure 4: Numbers of astrocytes, mature neurons and immature cells in the paralaminar nucleus of the monkey amygdala at different ages during postnatal development. Note that the total number of neurons (mature + immature) remains constant throughout postnatal development. N: Newborns; 3M: 3-month-old; 6M: 6-month-old; 9M: 9-month-old; 1Y: 1 year; 5-9Y: 5-9-year.

Figure 5: Numbers of oligodendrocytes in the six main nuclei of the monkey amygdala at different ages during early postnatal development. **A**: Lateral. **B**: Basal. **C**: Accessory basal. **D**: Paralaminar. **E**: Central. **F**: Medial. Error bars: \pm SD. N: Newborns; 3M: 3-month-old; 6M: 6-month-old; 9M: 9-month-old; 1Y: 1 year; 5-9Y: 5-9-year.

Figure 6: Neuronal soma size in the five main nuclei of the monkey amygdala at different ages during early postnatal development. **A₁-A₆:** Lateral. **B₁-B₆:** Basal. **C₁-C₆:** Accessory basal. **D₁-D₆:** Central. **E₁-E₆:** Medial.

Figure 7: Neuronal soma size in the paralamina nucleus of the monkey amygdala at different ages during early postnatal development. **A:** Newborns. **B:** 3-month-olds. **C:** 6-month-olds. **D:** 9-month-olds. **E:** 1-year-olds. **F:** 5-9-year-olds.

Figure 8: Relationship between oligodendrocyte number and volume of the main amygdala nuclei of monkeys from three months to one year of age and at 5-9 years of age. **A:** Lateral (volume = $32.747 + (0.0000053 \times \text{oligodendrocyte number})$; $R^2=0.23$; $F_{1,18} = 5.636$; $P = 0.029$). **B:** Basal (volume = $33.679 + (0.000011 \times \text{oligodendrocyte number})$; $R^2=0.49$; $F_{1,18} = 17.523$; $P < 0.001$). **C:** Accessory basal (volume = $19.259 + (0.0000092 \times \text{oligodendrocyte number})$; $R^2=0.42$; $F_{1,18} = 13.129$; $P = 0.002$). **D:** Paralamina (volume = $3.782 + (0.0000482 \times \text{oligodendrocyte number})$; $R^2=0.65$; $F_{1,18} = 34.620$; $P < 0.001$). **E:** Central (volume = $2.904 + (0.0000232 \times \text{oligodendrocyte number})$; $R^2=0.81$; $F_{1,18} = 76.982$; $P < 0.001$). **F:** Medial (volume = $4.548 + (0.00000686 \times \text{oligodendrocyte number})$; $R^2=0.18$; $F_{1,18} = 4.169$; $P = 0.056$).

Figure 9: Schematic representation of the postnatal development of the monkey amygdala circuits. L: lateral; B: basal; AB: accessory basal; C: central; M: medial; PL: paralamina. Black arrow: unmyelinated axons; black and white arrow: myelinated axons.

Table 1: Parameters used for the stereological analysis of the postnatal development of the monkey amygdala.

Table 2: Volume, cell numbers and neuronal soma size in the main nuclei of the monkey amygdala at different ages during early postnatal development.

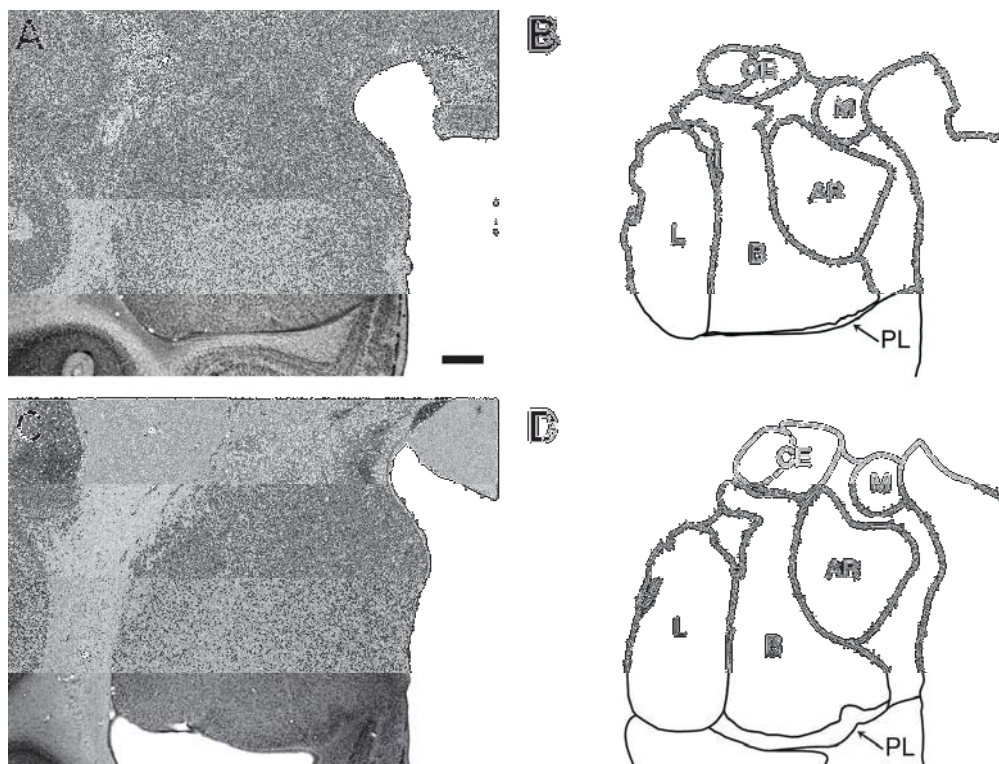
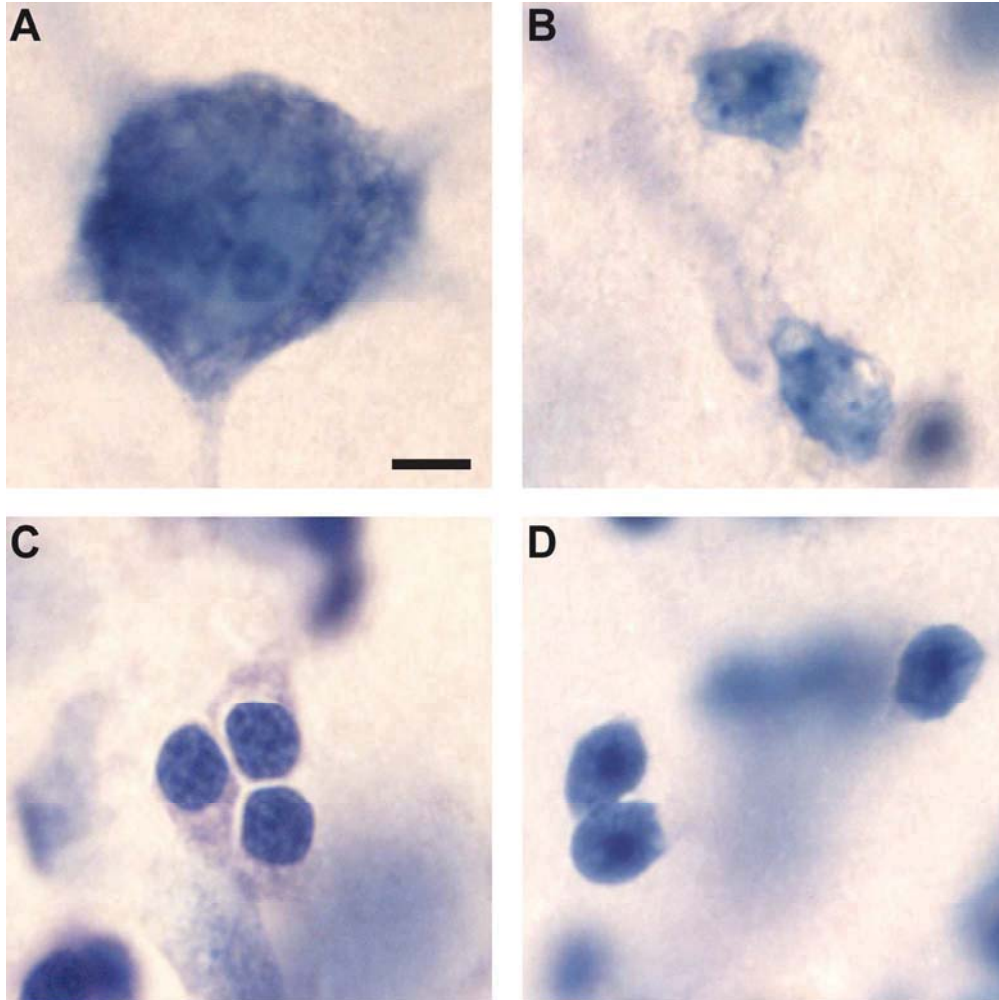


Figure 1: Low-magnification photomicrographs of a representative coronal section at a mid-rostrocaudal level of the rhesus monkey (*Macaca mulatta*) amygdala, which illustrate the locations of the six main amygdala nuclei in a newborn (A and B) and a 9-year-old (C and D) monkey. L: Lateral; B: Basal; PL: Paralaminar; AB: Accessory basal; CE: Central; M: Medial. Non-labeled areas include the remaining nuclei of the amygdala (see (Chareyron et al., 2011)). Scale bar in A: 1 mm, applies to all panels.
172x130mm (300 x 300 DPI)



171x171mm (300 x 300 DPI)

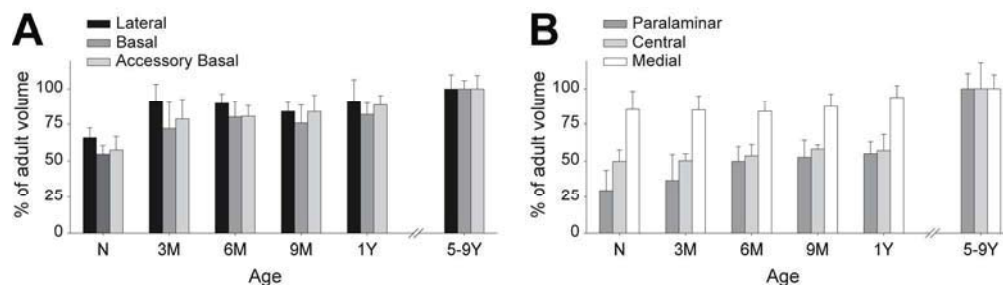


Figure 3: Volumes of the main nuclei of the monkey amygdala at different ages during early postnatal development (expressed as a percentage of the volume of the structure observed in 5-9-year-old monkeys: average \pm SD). A: Lateral, basal and accessory basal nuclei. B: Paralaminar, central and medial nuclei. N: Newborn; 3M: 3-month-olds; 6M: 6-month-olds; 9M: 9-month-olds; 1Y: 1-year-olds; 5-9Y: 5-9-year-olds. 170x47mm (300 x 300 DPI)

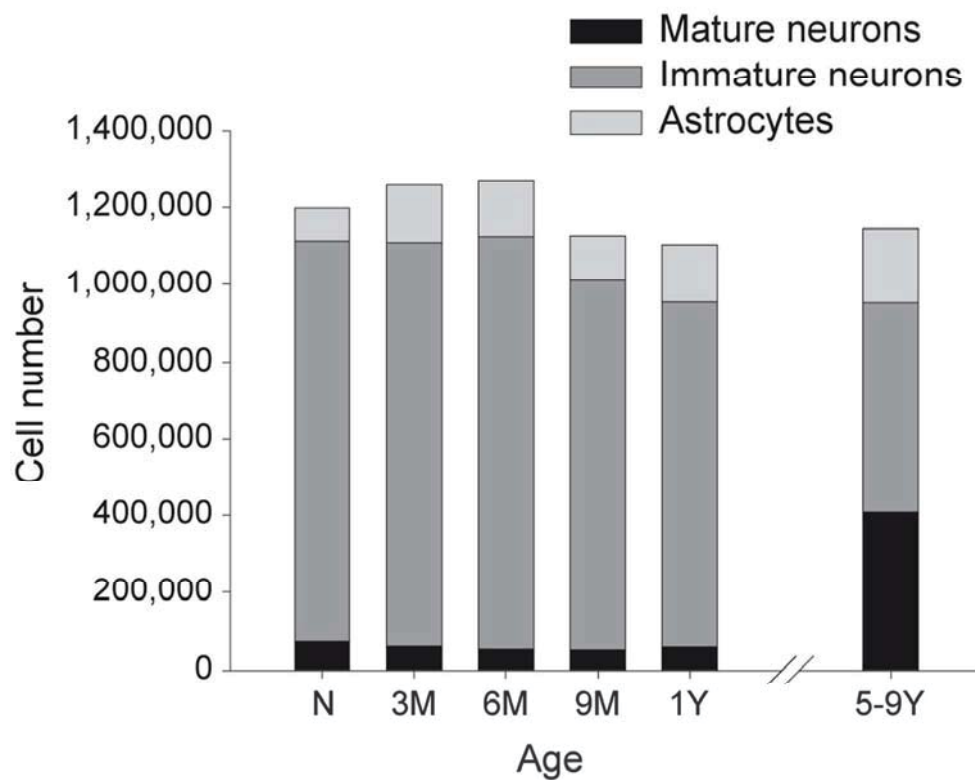


Figure 4: Numbers of astrocytes, mature neurons and immature cells in the paralaminar nucleus of the monkey amygdala at different ages during postnatal development. Note that the total number of neurons (mature + immature) remains constant throughout postnatal development. N: Newborns; 3M: 3-month-old; 6M: 6-month-old; 9M: 9-month-old; 1Y: 1 year; 5-9Y: 5-9-year.
85x68mm (300 x 300 DPI)

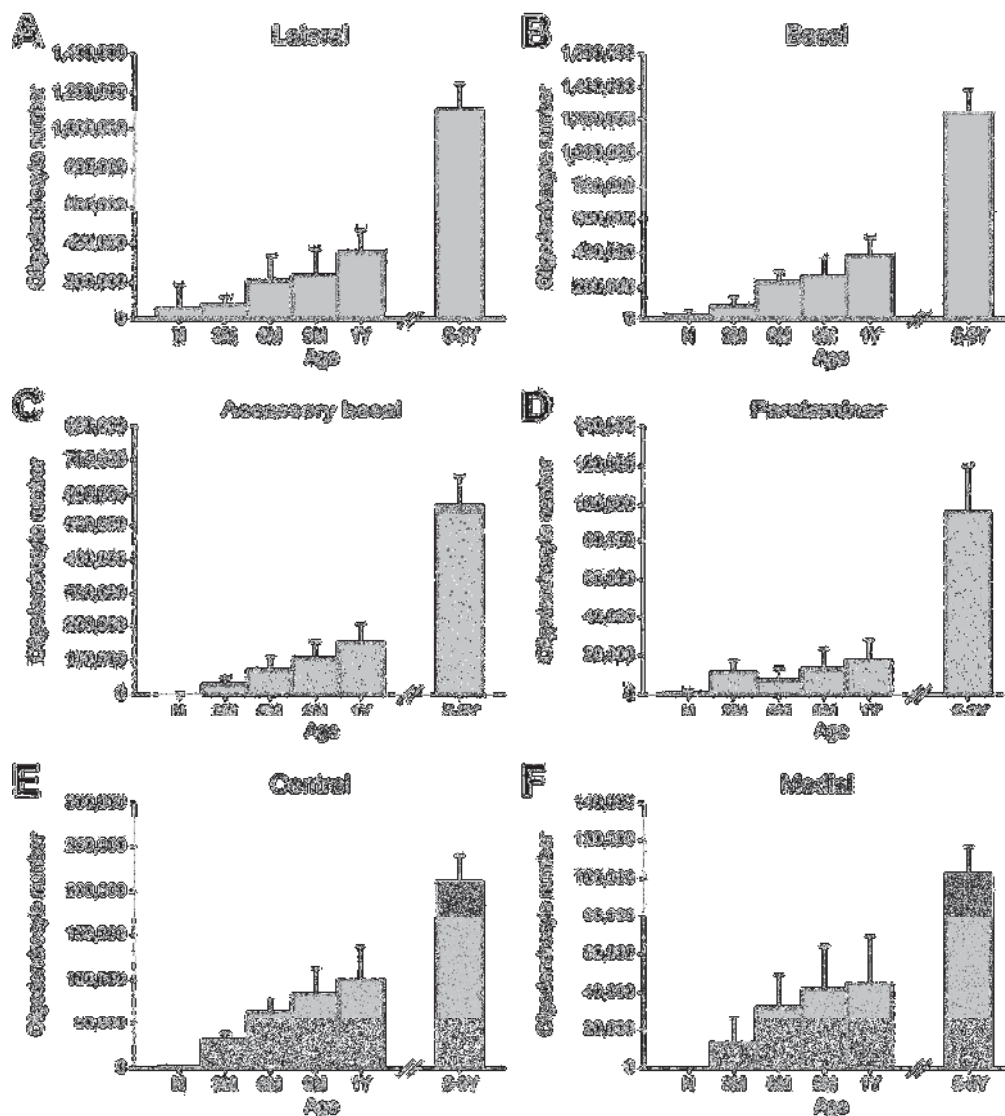


Figure 5: Numbers of oligodendrocytes in the six main nuclei of the monkey amygdala at different ages during early postnatal development. A: Lateral. B: Basal. C: Accessory basal. D: Paralaminae. E: Central. F: Medial. Error bars: +/- SD. N: Newborns; 3M: 3-month-old; 6M: 6-month-old; 9M: 9-month-old; 1Y: 1 year; 5-9Y: 5-9-year. 172x190mm (300 x 300 DPI)

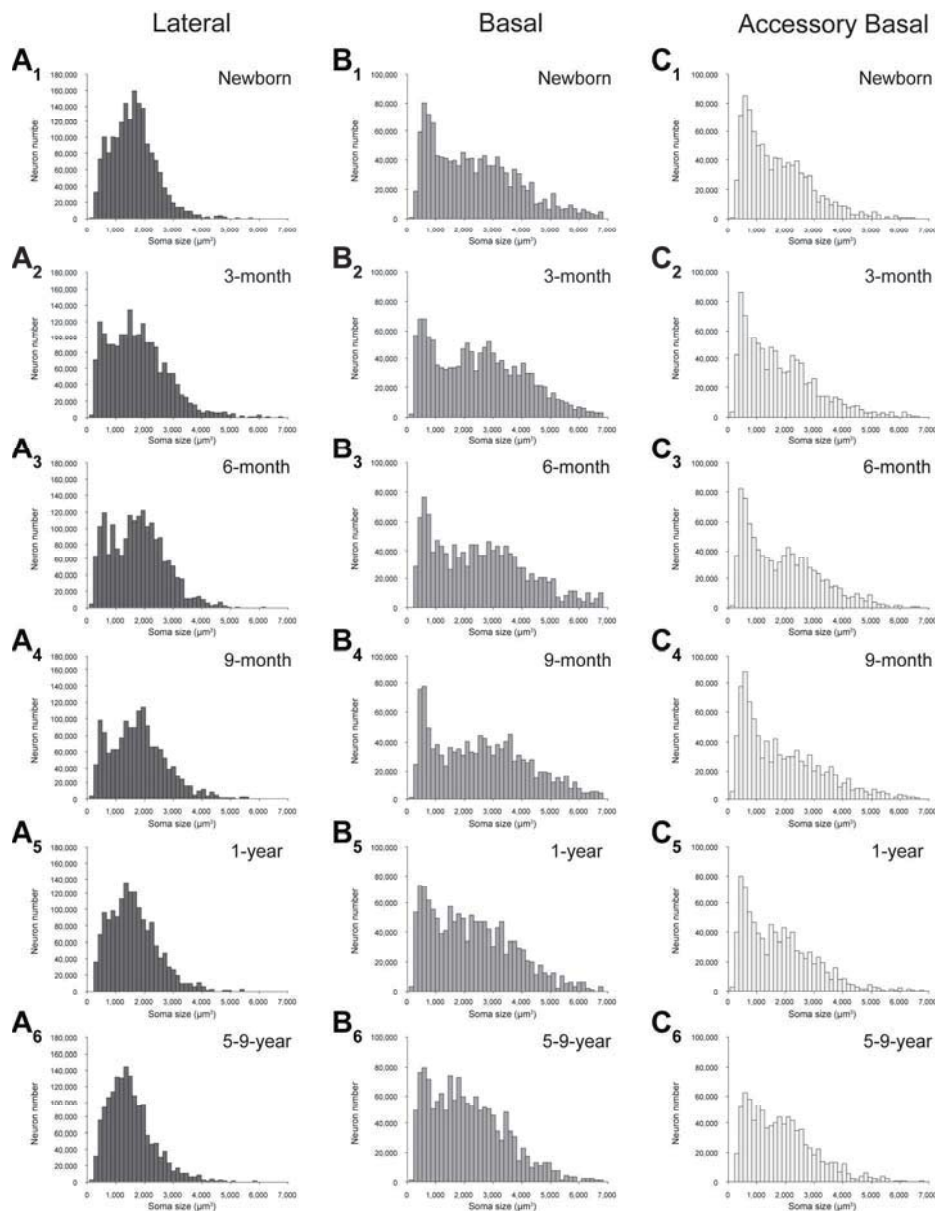


Figure 6: Neuronal soma size in the five main nuclei of the monkey amygdala at different ages during early postnatal development. A1-A6: Lateral. B1-B6: Basal. C1-C6: Accessory basal. D1-D6: Central. E1-E6: Medial.
177x230mm (300 x 300 DPI)

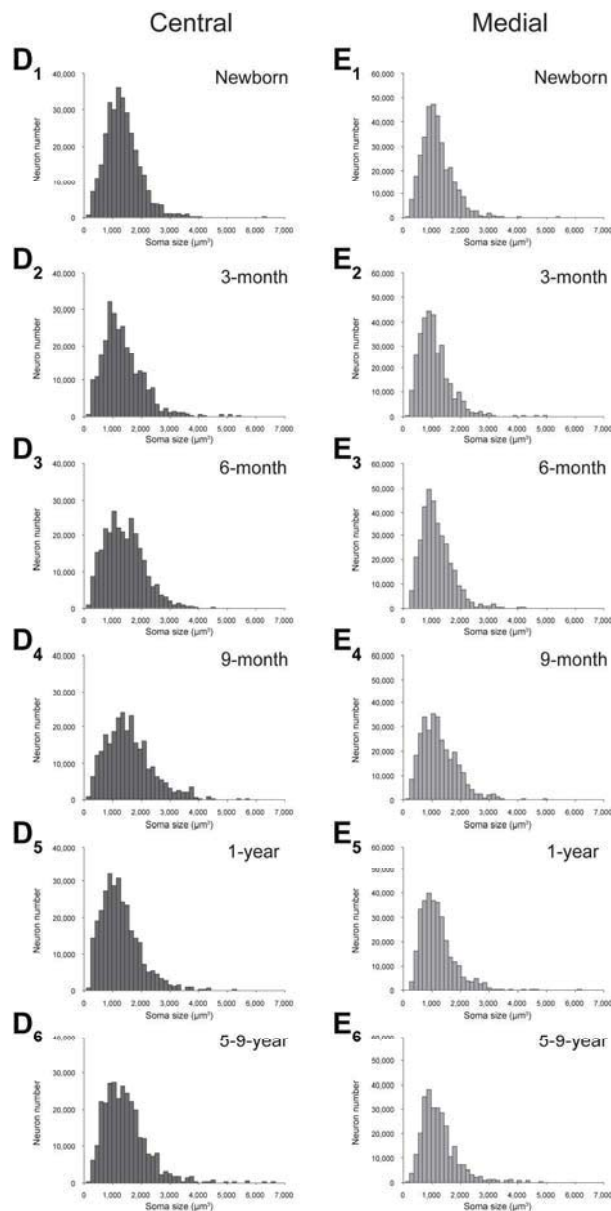


Figure 6: Neuronal soma size in the five main nuclei of the monkey amygdala at different ages during early postnatal development. A1-A6: Lateral. B1-B6: Basal. C1-C6: Accessory basal. D1-D6: Central. E1-E6: Medial.

114x230mm (300 x 300 DPI)

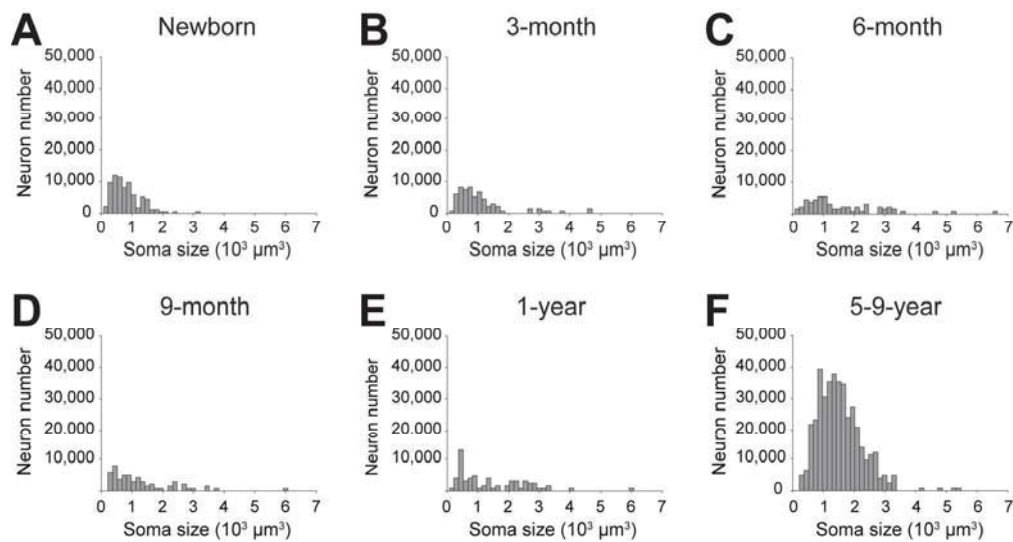


Figure 7: Neuronal soma size in the paralamina nucleus of the monkey amygdala at different ages during early postnatal development. A: Newborns. B: 3-month-olds. C: 6-month-olds. D: 9-month-olds. E: 1-year-olds. F: 5-9-year-olds.
172x91mm (300 x 300 DPI)

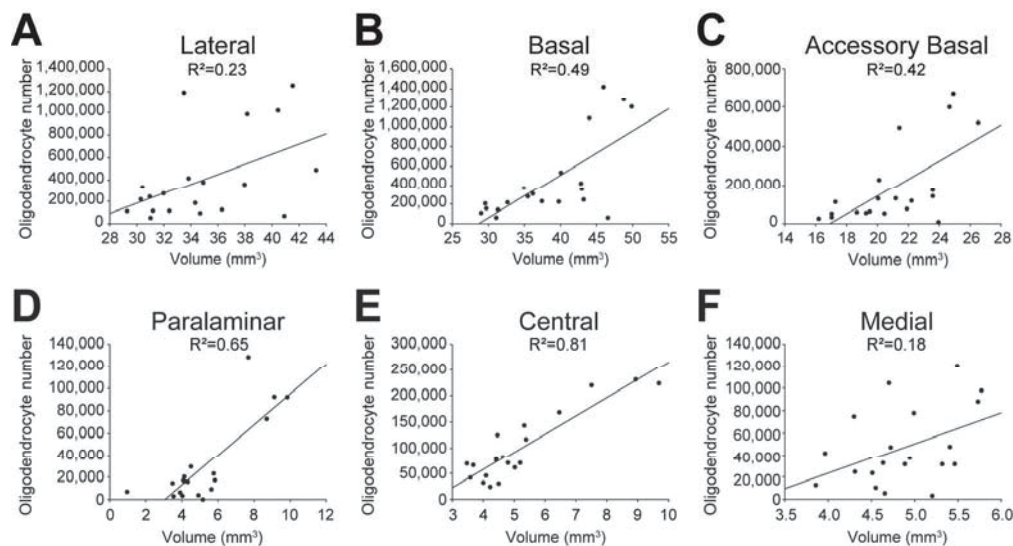


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170x92mm (300 x 300 DPI)

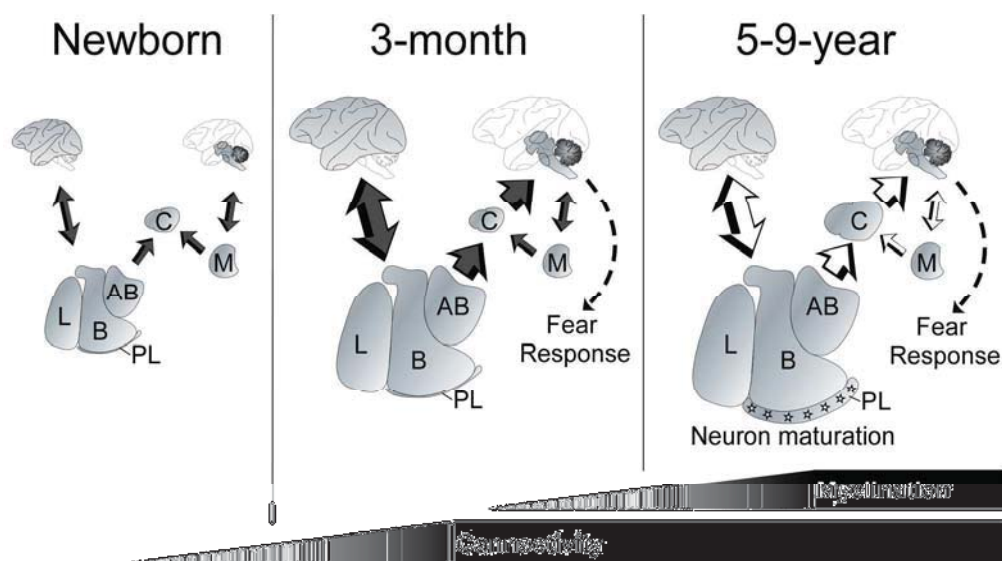


Figure 9: Schematic representation of the postnatal development of the monkey amygdala circuits. L: lateral; B: basal; AB: accessory basal; C: central; M: medial; PL: paralaminar. Black arrow: unmyelinated axons; black and white arrow: myelinated axons.
172x96mm (300 x 300 DPI)

Table 1: Parameters used for the stereological analysis of the postnatal development of the monkey amygdala.

Amygdala nucleus	Average number of sections (range)	Distance between sections (μm)	Scan grid (μm)*	Counting frame (μm)	Disector height (μm)	Guard zones (μm)	Average section thickness (μm)**	Neurons	Average number of cells counted (range)		
									Oligodendrocytes	Astrocytes	Immature neurons
Lateral	10 (8-13)	480	600 x 600	40 x 40	5	2	12.62 (8.22-15.97)	392 (257-653)	73 (0-280)	253 (180-439)	-
Basal	10 (9-12)	480	550 x 550	40 x 40	5	2	13.24 (8.78-16.74)	334 (232-536)	88 (0-351)	313 (207-449)	-
Paralaminar	11 (10-13)	480	250 x 250	20 x 20	5	2	11.91 (8.05-15.10)	39 (9-137)	8 (0-43)	46 (22-72)	322 (135-556)
Accessory basal	10 (8-12)	480	475 x 475	40 x 40	5	2	12.69 (8.98-15.32)	333 (248-522)	51 (0-216)	278 (208-419)	-
Central	7 (6-9)	480	300 x 300	40 x 40	5	2	12.42 (8.88-15.76)	266 (152-409)	71 (0-203)	255 (169-405)	-
Medial	8 (7-10)	480	375 x 375	40 x 40	5	2	11.78 (9.03-13.75)	197 (140-307)	23 (0-79)	161 (97-245)	-

* Scan grid was placed in random orientation. **Section thickness was measured at every other counting site.

Table 2: Volume, cell numbers and neuronal soma size in the main nuclei of the monkey amygdala at different ages during early postnatal development.

	Lateral			Basal			Accessory Basal			Paralaminar			Central			Medial		
NUCLEI VOLUME (mm ³)																		
Newborn	25.32	±	2.62	25.79	±	2.82	13.98	±	2.29	2.64	±	1.19	4.04	±	0.64	4.61	±	0.72
3-month	34.86	±	4.72	34.08	±	8.37	19.11	±	3.47	3.27	±	1.54	4.08	±	0.39	4.59	±	0.55
6-month	34.26	±	2.75	37.62	±	5.08	19.55	±	1.81	4.38	±	0.90	4.35	±	0.64	4.53	±	0.40
9-month	32.10	±	2.39	35.63	±	5.80	20.36	±	2.89	4.64	±	1.02	4.73	±	0.24	4.72	±	0.50
1-year	34.73	±	6.08	38.36	±	3.85	21.48	±	1.72	4.85	±	0.72	4.65	±	0.90	5.09	±	0.44
5-9-year	38.40	±	3.57	47.15	±	2.67	24.38	±	2.14	8.84	±	0.90	8.15	±	1.44	5.42	±	0.50
NEURON NUMBER																		
Newborn	1,770,271	±	163,872	1,232,759	±	220,811	951,634	±	196,808	75,024	±	15,929	312,095	±	22,590	343,439	±	32,907
3-month	1,925,343	±	123,564	1,328,170	±	112,143	959,225	±	56,020	61,922	±	11,089	274,211	±	23,266	322,184	±	42,384
6-month	1,865,763	±	341,660	1,296,522	±	110,063	939,943	±	64,479	55,446	±	32,295	280,798	±	27,801	348,810	±	48,376
9-month	1,550,860	±	113,446	1,227,670	±	159,012	969,833	±	62,240	53,343	±	36,445	269,968	±	29,756	320,119	±	63,193
1-year	1,598,605	±	284,359	1,347,395	±	129,287	881,131	±	71,369	60,640	±	5,811	300,997	±	69,074	316,043	±	9,063
5-9-year	1,592,284	±	128,381	1,445,882	±	138,651	885,352	±	101,469	408,051	±	49,453	297,079	±	30,889	282,622	±	25,053
ASTROCYTE NUMBER																		
Newborn	1,075,160	±	199,787	1,147,166	±	88,636	811,074	±	113,488	83,872	±	27,587	294,217	±	29,384	284,953	±	28,999
3-month	1,348,781	±	177,464	1,280,084	±	205,464	825,492	±	58,221	147,255	±	35,000	253,739	±	31,139	272,562	±	44,068
6-month	1,150,751	±	154,674	1,202,133	±	242,384	720,950	±	60,125	142,127	±	26,101	265,206	±	41,489	246,904	±	35,274
9-month	1,029,207	±	15,720	1,187,168	±	195,565	779,742	±	85,046	116,953	±	41,715	280,819	±	50,842	249,895	±	77,082
1-year	1,049,401	±	198,033	1,328,610	±	210,249	804,816	±	12,100	150,934	±	26,169	295,778	±	50,687	291,903	±	45,238
5-9-year	1,044,198	±	64,819	1,237,921	±	86,478	701,739	±	23,552	191,762	±	28,505	276,106	±	15,546	235,419	±	17,978
OLIGODENDROCYTE NUMBER																		
Newborn	65,897	±	116,192	18,973	±	37,946	1,668	±	2,013	602	±	1,205	815	±	941	0	±	0
3-month	88,714	±	34,991	94,902	±	45,779	29,149	±	19,368	11,492	±	5,919	31,473	±	7,393	13,323	±	13,275
6-month	208,438	±	128,030	228,086	±	68,389	72,778	±	37,834	7,644	±	6,015	64,864	±	14,405	32,680	±	16,572
9-month	237,718	±	129,714	267,359	±	104,294	103,735	±	50,864	13,359	±	9,665	84,303	±	28,442	41,880	±	22,442
1-year	355,195	±	114,080	384,490	±	107,272	153,342	±	51,464	17,538	±	10,532	100,881	±	35,533	45,027	±	24,144
5-9-year	1,112,480	±	131,622	1,247,140	±	139,668	572,196	±	82,235	96,422	±	23,397	210,656	±	28,544	102,661	±	13,889
NEURONAL SOMA VOLUME AVERAGE (μm ³)																		
Newborn	1,536	±	95	2,419	±	226	1,646	±	131	740	±	124	1,265	±	110	1,089	±	152
3-month	1,675	±	375	2,492	±	513	1,691	±	376	1,005	±	219	1,233	±	247	1,015	±	80
6-month	1,694	±	177	2,555	±	354	1,826	±	124	1,524	±	263	1,328	±	122	1,067	±	101
9-month	1,679	±	205	2,567	±	333	1,822	±	189	1,058	±	418	1,483	±	200	1,191	±	195
1-year	1,533	±	112	2,268	±	209	1,707	±	97	1,385	±	489	1,194	±	117	1,129	±	61
5-9-year	1,433	±	158	2,073	±	99	1,804	±	92	1,470	±	87	1,374	±	69	1,176	±	27
% of NUCLEUS VOLUME OCCUPIED BY NEURONAL SOMAS																		
Newborn	10.77	±	0.93	11.49	±	1.24	11.25	±	1.86	3.08	±	2.85	9.85	±	1.01	8.21	±	1.53
3-month	9.18	±	1.14	9.74	±	0.35	8.44	±	0.52	2.49	±	1.63	8.31	±	1.70	7.27	±	1.78
6-month	9.19	±	1.51	8.83	±	1.05	8.85	±	1.25	2.01	±	1.18	8.70	±	1.62	8.38	±	2.15
9-month	8.11	±	0.97	8.86	±	0.91	8.71	±	0.50	1.22	±	0.99	8.52	±	1.73	7.92	±	0.72
1-year	7.05	±	0.40	7.96	±	0.63	7.01	±	0.52	1.72	±	0.57	7.67	±	0.60	7.04	±	0.65
5-9-year	5.96	±	0.79	6.34	±	0.32	6.54	±	0.41	6.80	±	0.70	5.06	±	0.40	6.16	±	0.63

	Amygdala			Brain*		
VOLUME (mm ³)						
Newborn	116.34	±	15.24	44,655	±	4,338
3-month	158.10	±	25.97	55,767	±	3,585
6-month	156.12	±	12.00	56,857	±	5,511
9-month	159.84	±	16.67	53,475	±	5,541
1-year	166.26	±	10.11	56,188	±	5,910
5-9-year	193.86	±	12.86	52,360	±	4,875

*Refers to the volume of the telencephalon and diencephalon bilaterally.

Paralaminar			
IMMATURE NEURON NUMBER			
Newborn	1,038,954	±	247,977
3-month	1,046,627	±	290,699
6-month	1,069,391	±	188,256
9-month	956,429	±	208,996
1-year	893,367	±	71,509
5-9-year	544,588	±	78,783